

Corrigendum

" γ Sorbitol as a diluent in tablets"

In Drug Development and Industrial Pharmacy 11, 551-564, some mistakes were introduced in the text ; so, would you please read the new pages 558-559 in place of the old ones.

558 According to the above mentioned litterature, it could be thought that the 86-87° C melting form is the α form and the 91.7° C melting form, the β form.

Each sort of Sorbitol has its own use in food or pharmaceutical industries. Regarding their use in their solid state as diluents in tablet formulation, it seems that the more stable form is the γ one.

We tested it for hygroscopicity and we compared it with the following :

S_1 ($\alpha + \beta + \gamma$ Sorbitol)
 S_8 (much $\alpha + \beta$)
 S_{11} (very impure γ form)

After one month's storage at different relative humidities, we measured the water content increase of these four samples by the Karl Fischer method.

The results are displayed in table II.

TABLE II : The water content of different samples of Sorbitol after one month's storage at different relative humidities and at 20° C

| Relative humidity | Time 0 | After one month | | | |
|--------------------------------------|--------|-----------------|--------|--------|---------|
| | | 22% RH | 54% RH | 70% RH | 80% RH |
| Type of sorbitol : | | | | | |
| S ₂ (γ) | 0.40 % | 0.10 % | 1.20 % | 3.10 % | 17.90 % |
| S ₁ (α+β+γ) | 0.67 % | 1.30 % | 2.40 % | 4.60 % | 27.85 % |
| S ₈ (much α+β) | 0.25 % | 0.20 % | 0.50 % | 4.80 % | 21.00 % |
| S ₁₁ (very impure γ form) | 0.80 % | 0.85 % | 1.30 % | 4.75 % | 24.90 % |

Considering these results, we selected a pure γ Sorbitol for tablet formulation study.

II. Compression and tablet study of tablets prepared with γ Sorbitol

II.1. Compression study

TABLE III : Compression parameters of Sorbitol and Lactose tablets

x = upper punch displacement (volume of compression chamber remaining constant : depth = 1 cm)
 y_1 = maximum upper punch stress (KN)
 y_2 = maximum lower punch stress (KN)
 y_2/y_1 = is indicative of stress transmission through the powder during the compression

| Drug | Disintegrant | Diluant | x | $\frac{y_1}{m}$ | C.V. | $\frac{y_2}{m}$ | C.V. | y_2/y_1 |
|---------------|--------------|----------|-----|-----------------|------|-----------------|------|-----------|
| Aspirin | Maïze | Sorbitol | 408 | 10.4 | 2.1% | 7.8 | 2.3% | 0.75 |
| | Starch | Lactose | 418 | 9.8 | 2.0% | 7.9 | 2.2% | 0.81 |
| | Kollidon | Sorbitol | 324 | 5.2 | 4.8% | 4.2 | 5.0% | 0.81 |
| | CL | Lactose | 435 | 5.6 | 5.0% | 4.3 | 4.0% | 0.78 |
| | Ac Di Sol | Sorbitol | 403 | 5.8 | 5.4% | | | |
| | | Lactose | 403 | 5.4 | 4.4% | | | |
| Ascorbic Acid | Maïze | Sorbitol | 352 | 10.7 | 3.5% | 9.1 | 3.4% | 0.85 |
| | Starch | Lactose | 364 | 9.9 | 1.5% | 8.6 | 1.4% | 0.87 |
| | Kollidon | Sorbitol | 456 | 9.1 | 3.2% | 7.6 | 3.2% | 0.83 |
| | CL | Lactose | 487 | 11.1 | 2.8% | 9.2 | 2.8% | 0.84 |
| | Ac Di Sol | Sorbitol | 390 | 12.0 | 1.8% | | | |
| | | Lactose | 403 | 5.8 | 5.4% | | | |

Sorbitol/Lactose comparison : Except for the Ascorbic Acid/Ac Di Sol tablets, the results are nearly the same but for a sound discussion they must be compared with hardness values of resulting tablets.

II.2. Hardness

TABLE IV : Hardness of Sorbitol and Lactose tablets (in KN)

| | | | Hardness | | Hardness |
|---------------|-----------|----------|----------|-------|----------|
| | | | m | C.V. | y_1 |
| Aspirin | Maïze | Sorbitol | 124 | 7.7% | 1.19 |
| | Starch | Lactose | 90 | 10.8% | 0.91 |
| | Kollidon | Sorbitol | 64 | 11 % | 1.22 |
| | CL | Lactose | 60 | 11.8% | 1.07 |
| | Ac Di Sol | Sorbitol | 87 | 11.3% | 1.51 |
| | | Lactose | 50 | 11.8% | 0.92 |
| Ascorbic Acid | Maïze | Sorbitol | 54 | 8.3% | 0.51 |
| | Starch | Lactose | 29 | 8.3% | 0.29 |
| | Kollidon | Sorbitol | 65 | 10.7% | 0.71 |
| | CL | Lactose | 56 | 11.6% | 0.51 |
| | Ac Di Sol | Sorbitol | 91 | 8 % | 0.76 |
| | | Lactose | 52 | 12.9% | 0.44 |